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May 7, 2003, 15:14:54; Search time 134.5 Seconds (without alignments) 6443.584 Million cell updates/sec
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GenCore version 5.1.4_p5_4578
Copyright (c) 1993 - 2003 Compugen Ltd.

    protein search, using frame_plus_n2p model

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Database

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0 Maximum DB seq length: .2000000000

ALIGNMENTS

RESULT 1 AAY22426

AAY22426 standard; Protein; 1083 AA

AAY22426;

28-SEP-1999 (first entry)

Human brain specific potassium channel protein sequence.

Brain specific potassium channel; human; central nervous system disorder; dementia; cerebral ischaemic sclerosis; therapy.

WO9937677-A1.

Homo sapiens.

29-JUL-1999

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98US-0091469. 99US-0116621.

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comprising an alpha subunit of the voltage-gated potassium channel comprising an alpha subunit of the voltage-gated potassium channel comprising an alpha subunit of the voltage-gated potassium channel seq (vGPCs). It is a member of the Kv (Voltage gated potassium superfamily, gated ether a go-go) family and Elk subfamily of potassium channel seq (ether a go-go) family and Elk subfamily of potassium channel commonners. helk gene is mapped to chromosome 12qi3. It is isolated from brain and maintains the resting potential and controls excitability of the cell. It has antimigrane, cerebroprotective, antipsychotic, of the cell. It has antimigrane, cerebroprotective, antipsychotic, concurprotective and anticonvulsant activity. The helk polypeptide can be used to screen for modulators of VGPCs, that are useful for treating used to screen for modulators of VGPCs, that are useful for treating abnormal ion flux disorders. CNS disorders such as migraines, hearing and viston problems, setaures, psychotic disorders and to prevent atroves. It can be used as a marker for diagnosis of diseases linked to this gene and also as reporter molecule in detection systems. The this gene and also as reporter molecule in detection systems. The polynucleotide is useful for gene therapy, to rectify ELK expression.
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                                                                                 Wickenden A;
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Voltage gated potassium channel subunit; VGPCs; hElk; human; stroke; Rv superfamily; Eag family; ether a go-go; Elk subfamily; modulator; chromosome 12q13; resting potential; cell excitability; selzure; marker; CNS; migraine; treat; hearing/vision problem; psychotic; anticonvulsant; ion flux disorder; reporter molecule; detection; gene therapy; antimigrane; cerebroprotective; neuroprotective; antipsychotic.
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Mismatches:
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ArgLysSerGlyLeuProPheTrpCysLeuLeuAspValIleProIleLysAsnGluLys
                    GGGGAGGTGGCTCTCTTCCTAGTCTCTCACAAGGACATCAGCGAAAACCAAGAACCGAGGG
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Location/Qualifiers

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The present sequence is the human ELK (hElk) polypeptide variant #2, comprising an alpha subunit of the voltage-gated potassium channel comprising an alpha subunit of the voltage gated potassium superfamily, (VGPCS). It is a member of the Kv (Voltage gated potassium superfamily, ce (with a go-go) family and Elk subfamily of potassium channel components. BENG gene is mapped to chromosome 12q13. It is isolated monomers. BENG gene is mapped to chromosome 12q13. It is isolated from brain and maintains the resting potential and controls excitability of the cell. It has antimigrane, cerebroprotective, antipsychotic, of the cell. It has antimigrane, cerebroprotective, antipsychotic, cof the cell. It has antimigrane, cerebroprotective, antipsychotic, cof the cell. It has antimigrane, cerebroprotective, antipsychotic, cof the nor flux disorders, CNS disorders such as migraines, hearing abnormal ion flux disorders, CNS disorders and to prevent strokes. It can be used as a marker for diagnosis of diseases linked to strokes. It can be used as a marker for diagnosis of diseases linked to chinucleotide is useful for gene therapy, to rectify ELK expression. CNP polynucleotide is useful for gene therapy, to rectify ELK expression. CNP of the present sequence is not found in the specification but the present manner of the sequence found in page 62.
                                                                                                                                                                                                                                                                                                                                                                                                                                                     Novel polynucleotides and polypeptides of human ELK, a voltage-gated potassium channel subunit useful for treating ELK mis-expression and to screen for inhibitors and activators of such channels
                            amino acids'
                                               /label= Extended_P-S6_region // note= "Extended Pore-S6 region with conserved acids"
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                label- P-S6_region
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Conservative:
Mismatches:
Indels:
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421 Asn	<pre>Qy 1321 GAGCTGGGGGGCCGTCGCTGCGCAGCGCCTACATCACCTCCCTC</pre>	1381 AGCAGCCTCACCAGGGGCTTCGGCAACGTGTCCGCCAACGGGACACGGGAAAGATC 14	461 SerSerLeuThrSerValGlyPheGlyAsnValSerAlaAsnThrAspThrGluLys1le 48	VY 1441 TICICCATCTGCACCATGGTCATGGGCGCCCTGATGCACGCGGTGTTTTGGGAACGTG 1500 Whill	c 15	501 ThralaileileGinArgMetTyrAlaArgArgPheLeuTyrHisSerArgThrArgAsp 520	Qy 1561 CTGCGCGACTACATCCGCATCCACCGTATCCCCAAGCCCCTCAAGCAGCGCATGCTGGAG 1620	0.4° UTSPECTORECONSTRUCTOR OF THE CONTRACT OF	Db 541 TyrPheGlnalaThrTrpAlaValAsnAsnGly1leAspThrThrGluLeuLeuGlnSer 560	Qy 1681 CTCCCTGACGACGACGACAACATCGCCATGCACCTGCAAGGAGGTCCTGCAGGG 1111 1111 1111 1111 1111 1111 1111 1111 1111 1111 1111 1111 1111 1111 1111 1111 111 1111 <td< td=""><td>1741 CCACTGTTTGAGGGGGGCCAGCCGGGGGGGGCTGCCGGGGCACTGTCTGGCGCCCTGGGGCCC 18</td><td>581 ProLeuPheGluAlaAlaSerArgGlyCysLeuArgAlaLeuSerLeuAlaLeuArgPro 600</td><td>1801</td><td>1861 TITGICIGCTCTGGCTCCATGGAGGTGCTCAAGGGTGGCACCGTGCTCGCCATCCTAGGG 19</td><td>1921 AAGGGCGACTTCARTCCTTTTTTTTTTTTTTTTTTTTTTT</td><td>64</td><td>1981 GACGTGAAGGGCTGACGTACTGCGTCTGCAGTGTCTGCAGCTGGCTG</td><td>2041 ACCUMEDO</td><td>681 Serie</td><td>2101 AGCTACAACCTGGGTGCTCGGGGGGGCTCTGCAGACGTGGACACCAGCTCCCTGAGCGGC 2160 </td><td>2161 GACAATACCCTTATGTCCACGCTGGAGGAGAGAGAGAGAG</td><td>2221 GTCTCCCCAGCCCCAGCTGATGAGCCCTCCAGCCCCCTGCTGCCCTGCTGCCTGC</td><td>CCCGCCTCGTCTAGGT 23</td><td>CCTCTGCTCCC 24</td><td></td></td<>	1741 CCACTGTTTGAGGGGGGCCAGCCGGGGGGGGCTGCCGGGGCACTGTCTGGCGCCCTGGGGCCC 18	581 ProLeuPheGluAlaAlaSerArgGlyCysLeuArgAlaLeuSerLeuAlaLeuArgPro 600	1801	1861 TITGICIGCTCTGGCTCCATGGAGGTGCTCAAGGGTGGCACCGTGCTCGCCATCCTAGGG 19	1921 AAGGGCGACTTCARTCCTTTTTTTTTTTTTTTTTTTTTTT	64	1981 GACGTGAAGGGCTGACGTACTGCGTCTGCAGTGTCTGCAGCTGGCTG	2041 ACCUMEDO	681 Serie	2101 AGCTACAACCTGGGTGCTCGGGGGGGCTCTGCAGACGTGGACACCAGCTCCCTGAGCGGC 2160	2161 GACAATACCCTTATGTCCACGCTGGAGGAGAGAGAGAGAG	2221 GTCTCCCCAGCCCCAGCTGATGAGCCCTCCAGCCCCCTGCTGCCCTGCTGCCTGC	CCCGCCTCGTCTAGGT 23	CCTCTGCTCCC 24	
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epilepsy; migraine; cell proliferation disorder; cancer;
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                                                                                                                                                                                                                       the present sequence represents a human potassium channel protein of the ERG family, which is designated Herg4 (human erg related gene 4). The Herg4 polypeptides and polymucleotides are useful in the treatment of epilepsy, migraine, cell proliferation disorders, cancer, comportemental troubles, and to prevent or alter the effect of endogenous neurotransmitters and hormones. Antibodies against Herg4 are also useful for the treatment of cerebral, cardiac and renal ischemias, brain and cardiac diseases, inflammation, pain, and to mimic or antagonize the effect of endogenous neurotransmitters and hormones.
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  trouble; neurotransmitter; hormone; ischemia; cardiac disease; inflammation; pain.
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cell proliferation
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Matches:
Conservative:
Mismatches:
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for treatment of epilepsy, migraine,
                                                                                                                                                                                                           claim 11; Page 45-48; 48pp; English.
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Best Local Similarity:
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brain disease;
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                                                              481 AAAGGCTTCAATGCCAACCGGCGGCGGAGCCGGGCCGTGCTCTACCACCTGTCCGGGCAC
GlyGluValAlaLeuPheLeuValSerHisLysAspIleSerGluThrLysAsnArgGly
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Voltage gated potassium channel subunit; VGPCs; hElk; human; stroke; Kv superfamily; Eag family; ether a go-go; Elk subfamily; modulator; chromosome 12q13; resting potential; cell excitability; seizure; marker; CNS; migraine; treat; hearing/vision problem; psychotic; anticonvulsant; ion flux disorder; reporter molecule; detection; gene therapy; antimigrane; cerebroprotective; neuroprotective; antipsychotic.
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/label= P-S6_region
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rne present sequence is the number of the Voltage-gated potassium channel comparising an alpha subunit of the Voltage-gated potassium channel (VGPCS). It is a member of the KV (Voltage gated potassium channel sequences. It is a member of family and Elk subfamily of potassium channel monomers. hElk gene is mapped to chromosome 12q13. It is isolated monomers. hElk gene is mapped to chromosome 12q13. It is isolated from brain and maintains the resting potential and controls excitability of from brain and maintains the resting potential and controls excitability of the cell. It has antimigrane, cerebroprotective, antipsychotic, neuroprotective and anticonvulsant activity. The hElk polypeptide can be used to screen for modulators of VGPCS, that are useful for treating and vision problems, selzures, psychotic disorders and to prevent strokes. It can be used as a marker for disquosts of diseases linked to strokes. It can be useful for gene therapy, to rectify Elk expression. Note: The present sequence is not found in the specification but child for the helk amino acid sequence found in page 62.
                                                                                                                                                                                                                                                                                                                                                                                                                      Novel polynucleotides and polypeptides of human ELK, a voltage-gated potassium channel subunit useful for treating ELK mis-expression and to screen for inhibitors and activators of such channels
/label- Extended_P-S6_region /note- "Extended Pore-S6 region with conserved amino acids"
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              present sequence is the human ELK (hElk) polypeptide variant
                                                                             type Ala substitued with Ser'
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CCCGTGGTCTACTGCTCTGATGCTTCTGTGACCTCACGGGCTTCTCCCGGGCTGAGGTC 180 240 241 CAACAGATCCGCAAGGCCCTGGACGAGCACAAGGAGTTCAAGGCTGAGCTGATCCTGTAC 300 120 20 40 61 CGCTTCGACGGCACGCACAGTAACTTCGTGCTGGGCAACGCCCAGGTGGCGGGGGCTCTTC 11 ArgPheAspGlyThrHisSerAsnPheValLeuGlyAsnAlaGlnValAlaGlyLeuPhe ATGCCGGCCCATGCGGGGCCTCCTGGCGCCTCAGAACACCTTCCTGGACACCATCGCTACG 1083 1081 US-09-965-830-1_COPY_6_3257 (1-3252) x AAY44781 (1-1083) Matches: Conservative: Mismatches: Indels: 5695.00 99.918 99.828 93.538 Percent Similarity: Best Local Similarity: Alignment Scores: Sequence Best Local S Query Match .. Q 121 181 61 41 Score: Pred. g g

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ESK1; eag similar K+ channel; potassium channel associated disorder; neurological; Alzheimer's disease; anxiety; panic; autism; hyperactivity; obsessive-compulsive disorder; schizophrenia; Huntington's disease; epilepsy; cardiovascular; musculoskeletal; prolliferative, cancer; ESK channel blocker; nootropic; neuroprotective; antidepressant; tranquilizer; neuroleptic; antiParkinsonian; cardiant; cytostatic;
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        This represents a eag similar K+ channel (ESK) polypeptide (hESKI). The hESKI protein can be expressed by standard recombinant methodology. The ESK polypeptide, polynucleotides and antibodies are useful for treating and diagnosing various potassium channel associated disorders such as neurological disorders, e.g. Alzheimer's disease, depression, anxiety, panic, obsessive-compulsive disorders, attention deficit, epilepsy; panic, obsessive-compulsive disorders, attention deficit, epilepsy; hyperactivity disorders, autism, schizophrenia, Huntington's disease and hyperactivity disorders such as cancer. The ESK polynucleotide is also useful for synthesis of ESK and gene mapping. The polypeptide can be used in an assay to identify molecules such as synthetic diugs, antibodies, peptides or other molecules which have an effect on the activity of the ESK channel.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                Novel eag similar potassium channel polypeptide useful for treating various neurological, cardiovascular, musculoskeletal and proliferative
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The present sequence is a ERG-like protein 1 (ERG-LP1) which is a member of ERG potassium channel family. This sequence is from a full length clone jlkba25610 which was derived from monkey hippocampal library. ERG-LP1 is expressed exclusively in the brain. Highest expression is found in cortical regions, hippocampus, caudate and amygdala. The protein functions as a potassium channel modulator and has neuroprotective, antiParkinsonian, anticonvulsant, antidepressant,
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note= "Casein kinase II phosphorylation site"
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/note= "Protein kinase C phosphorylation site"
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te= "Casein kinase II phosphorylation site"
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      Monkey; potassium channel molecule; ERG-like protein 1; ERG-LP1; neuroprotective; antiParkinsonian; anticonvulsant; antidepressant; neuroleptic; nootropic; treatment; CNS disorder; central nervous system; potassium channel mediated disorder; epilepsy; Alzheimer's disease; Parkinson's; multiple sclerosis; depression; schizophrenia; amnesia.
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           Analor determinant of ion selectivity in potassium
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This sequence represents the human K+Hnov14 potassium channel.
K+Hnov proteins have a high degree of homology to known potassium channels and may be alpha subunits, which form the functional channel, or accessory subunits that act to modulate the channel activity.
K+Hnov14 is a 6 transmembrane domain, voltage gated potassium channel. The gene's chromosomal location is 12q14, determined via PCR chromosomal localisation using primers AAZ11928 and AAZ11929.
K+Hnov cDNAs were isolated by extension of expressed sequence tags (ESTS) which were related but not identical to known human potassium channels. Potential polymorphisms detected as sequence variants between multiple independent clones. Potassium channels have critical roles in various cell types and biochemical pathways. Defective potassium channels
GluProProAlaSerGlyAspLeuCysSerGluProSerThrProAlaSerProPro 1020
                                                                                                                                                      Potassium channel; ataxia; arrhythmia; epilepsy; Bartter's syndrome; cardiovascular disorder; CNS disorder; renal disorder.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       New nucleic acids encoding mammalian K+Hnov potassium channel proteins, useful for the diagnosis and treatment of episodic ataxia with myokymia, cardiac arrhythmia, epilepsy and Bartter's syndrome
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     a variant form of
                                                                                 GAGATGGTGCTTATTGGCTGCCATGGCTCTGGCACAGTCCAGTGGACCCAGGAAGAAGCC
                           TCTGAGGAAGGGGCTAGGACTGGGCCCGCAGACCTGTGAGCCAGGCTGAGGCTACCAGC
                                          /note= "Arg substituted by Cys in K+Hnov14"
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                                                                                                                                                                                                                                                                                       standard; Protein; 1082 AA
                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Location/Qualifiers
                                                                                                                                                                                                                                                                                                                                                                           Human potassium channel K+Hnov14
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98US-0076687.
98US-0095836.
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07-AUG-1998;
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are known to cause four human diseases: episodic ataxia with myokymia; cardiac arrhythmia (long Or syndrome); epilepsy; and Bartter's syndrome. As potassium channels are critical components of virtually all cells, it is likely that abnormal potassium channels are also implicated in certain renal, cardiovascular and central nervous system (CNS) disorders. Nucleotides encoding K+Hnov proteins may be used for them. They may be used to produce compositions that modulate the expression and function of the K+Hnov protein and in studying the blochemical pathways associated with it. They may also be used for the recombinant production of K+Hnov protein in fermentation cultures. Additionally, such nucleotides may be used in gene therapy protocols for the treatment of diseases associated with abnormal potassium channels.
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1082 AA; Sequence

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Alignment Sc Pred. No.: Score: Percent Simi Best Local S Query Match:	gnment d. No. re: cent t Loce ry Mat	<pre>lent Scores: No.: t Similarity: local Similarity: Match:</pre>	0 5649.50 99.358 99.788 20	Length: Matches: Conservative: Mismatches: Indels: Gaps:	1082 1076 0 6 6 1	
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οy	61		CGCACAGTAACTTCGT	GCTGGGCAACGCCCA		120
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qq	9		rillillillil 7sAlaCysSerPheLeu		MetGlnArgGlyCysAlaCysSerPheLeuTyrGlyProAspThrSerGluLeuValArg	79
ογ	241		AGGCCCTGGACGAGCAC	CAAGGAGTTCAAGGC		300
οqα	80				GInGln1leArgLysAlaLeuAspGluH1sLysGluPheLysAlaGluLeuIleLeuTyr 🤅	66
ολ	301		CCCGTTCTGGTGTCTC	CCTGGATGTGATACC		360
qq	100				ArgLysSerGlyLeuProPheTrpCysLeuLeuAspVallleProlleLysAsnGluLys	119
οy	361		CTTCCTAGTCTCTCAC	CAAGGACATCAGCGA	_	420
qq	120				GlyGluvalAlaLeuPheLeuValSerHisLysAspIleSerGluThrLysAsnArgGly	139
δλ	421		GAAGGAGACAGGTGG	TGGCCGGCGGATA		480
qq	140		PLYSGluThrGlyGly		GlyProAspArgTrpLysGluThrGlyGlyGlyArgArgArgTyrGlyArgAlaArgSer	159
δλ	481		CAACCGGCGGCGGAGC	CCGGGCCGTGCTCTA		540
qq	160		aAsnArgArgArgSer		LysGlyPheAsnAlaAsnArgArgArgArgAlaValLeuTyrHisLeuSerGlyHis	179
δ	541	CTGCAGAAGCAGCC	CAAGGGCAAGCACAAG	SCTCAATAAGGGGGT		009
QQ	180				Inninininininininininininininininininin	199
οχ	601		CAAAGTAGCCGCCATC	CGGAAGTCGCCCTT		099
മ	200		rLysValAlaAlaIle	argiysSerProPhe	AsnLeuProGluTyrLysValAlaAlaIleArgLysSerProPheIleLeuLeuHisCys 2	219
ογ	661	GGGCACTGAGAGC	CACCTGGGATGGCTTC	PATCCTGCTCGCCAC	ACTUTATETEGCTGTC 7	720
qa	220	GlyAlaLeuArgAl			GlyAlaLeuArgAlaThTTTpAspGlyPheIleLeuLeuAlaThrLeuTyrValAlaVal 2	39

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                                                    CGTACCACATTCGTGTCCAAGTCGGGCCAGGTGTTTGCCCCCAAAGTCCATTTGCCTC
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Human; potassium channel molecule; ERG-11ke protein 2; ERG-LP2; neuroprotective; antiParkinsonian; anticonvulsant; antidepressant; neuroleptic; nootropic; treatment; CNS disorder; central nervous system; potassium channel mediated disorder; epilepsy; Alzheimer's disease; Parkinson's; multiple sclerosis; depression; schizophrenia; amnesia; chromosome 3p21.3-24.3.
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/note= "Transmembrane region cyclic nucleotide gated
1000 GluProProAlaSerGlyAspLeuCysSerGluProSerThrProAlaSerProProPro
                                                                                GAGCCCCTGCCTCAGGAGACCTCTGCTCTGAGCCCCAGCACCCCTGCCTCCTCCT
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                                               "Protein kinase C phosphorylation site"
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159..161
/note= "Protein kinase C phosphorylation
216..218
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(MILL-) MILLENNIUM PHARM INC

Curtis RAJ;

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21-JUL-1999; 21-JUL-1998;

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/note= "Protein kinase C phosphorylation site" / 733..735 //note= "Protein kinase C phosphorylation site" / 733..735 //note= "Protein kinase C phosphorylation site" / 730.000 //note= | Protein kinase C phosphorylation site"
/note= "Protein Aimsec C phosphorylation site"
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/note= "Casein kinase II phosphorylation site"
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/note= "Casein kinase II phosphorylation site"
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/note= "Protein kinase C phosphorylation
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161..164
/note= "cAMP and cGMP dependent protein kinase
phosphorylation site"
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WO200005346-A1

03-FEB-2000

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The present sequence is a full-length ERG-like protein 2 (ERG-LP2) which is a member of ERG potassium channel family. This sequence is a result of additional sequencing of clone jlhbaa042h05 which was obtained from human brain library. ERG-LP2 gene is mapped to human chromosome 3p21.3-24.3, between markers WI-4218 and RP_LIS_I. It is predominantly expressed in the brain.

The protein functions as a potassium channel modulator and has neuroprotective, antiparkinsonian, anticonvulsant, antidepressant, neuroleptic and nootropic activities. The present sequence is useful for treating several potassium channel mediated disorders (CNS disorders) such as Alzheimer's disease, Parkinson's disease, multiple sclerosis, epilepsy, depression, schizophrenic disorders and amnesia.
                                                                molecule useful in treating as Alzheimer's disease, multiple
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      Length:
Matches:
Conservative:
Mismatches:
Indels:
                                                        Novel gene encoding potassium channel
central nervous system disorders such
sclerosis, and schizophrenia
                                                                                                                                       Example 1; Fig 8; 144pp; English.
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2529.50
60.24%
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WPI; 2000-182682/16.
N-PSDB; AAZ50455.
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AAAGGCTTCAATGCCAACCGGCGGAGCCGGGCCGTGCTCTACCACCTGTCGGG ::::::	CACCTGCAGAAGCAGCCCAAGGGCAAGCACAAGCTCAATAAGGGGGTGTTTGGGGAGAAA ::::::	CCAAACTIGCCTGAGTACAAAGTAGCCGCATCCGGAAGTCGCCCTTCATCCTGTTGCAC	IGTGGGGCACTGAGAGCCACCTGGGATGGCTTCATCCTGCTCGCCCACTCTATGTGGGT :::	GTCACTGTGCCCTACAGCGTGTGTGTGAGCACAGCACGGAGCCCAGTGCCGCCGCGGC	CCGCCCAGCGTCTGTGACCTGGCCGTGGAGGTCCTCTTCATCCTTGACATTGTGCTGAAT :::	TICCGTACCACATICGIGICCAAGICGGGCCAGGIGGIGITIGCCCCAAAGICCATITGC	CTCCACTACGTCACCACCACCTGGTTCCTGGTGGTCATCGCAGCGCTGCCCTTTGACCTG :::	CTACATGCCTTCAAGGTCAACGTGTACTTCGGGGCCCATCTGCTGAAGACGGTGCGCCTG	CJGCGCCJGCTGCCJGCTJCCGCGGCJGGACCGGJACTCGCAGJACAGCGCCGTGGJG 	CTGACACTGCTCATGGCCGTGTTCGCCCTGCTCCGCGCATGGCTCGCCTGCGTTTT	TACATTGGCCAGCGGGAGATCGAGAGCAGCAATCCGAGCTGCCTGAGATTGGCTGGC				CTCAGCAGCCTCACCAGCGTGGGCTTCGGCAACGTGTCCGCCAACACGGACACCGGAGAAG		
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Qy Db	1558	acccrcaaccaccargcrg 161
QY Db	QY 1618 GAGTACTTCCAGGCCACCTGGGCGGTGAACAATGGCATU	CAATGGCATCGACACCGAGCTGCTGCAG 1677 :: :: nAsnGlyIleAspSerAsnGluLeuLeuLys 528
oy Oy	1678	CGCCATGCACAGGAGGTCCTGCAG 1737
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VQ qq	1798 CCCG 569 ThrS	CCTTCTGCACGCCGGCGAGTACCTCATCCACCAGGCGATGCCCTGCAGGCCCTC 1857 ::
δy G	1858	TACTTIGICIGCTCTGGCTCCATGGAGGTGCTCAAGGGTGGCACCGTGCTCGCCATCCTA 1917
S d	1918	GCCCCGGCGGGAGCAGGTGGTAAAGGCCAAT 1977
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g G	: 8 689	:: erMetValSerGlnSerGluProLysGlyAsnGlyAsnIleAsnLysArgLeuProSer 708
oy d	2185	GAGGAGAAGGAGACAGATGGGGAGGGCCCCAGGGTCTCCCCAGCC 2232
δy	2233 CCAGCTGA	GCACCTCCTCATCCTCAGCT 2
a d	727	YSIIITAIGGTGGCAGAGGGAGG 23
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QY	2353 CCAGGCAGGGC	AGGCTGGCCCTTGCT2397
q	739 AsnLysLysVa	sniysalafyrLeuGlyLeuSerLeuLysGlnLeuAlaSer 758
δy	2397	23.
qq	759 GlyThrValProPheHisSerProlle	gvalSerArgSerAsnSerProLysThrLys 778
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2545 GAATGTAGCAGCAGCCCTCCCTGGA---CCAGAGAGGGGCCTGCTCACTGTTCCCCAT 2601
                                                                                                                      2602 GGGCCCAGCGAGGCAAGGAACACACACACACTGGACAAGCTTCGGCAGGCGGTGACAGAG 2661
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839 GluProArgileSerProProLeuGlyAspProGluileGlyAlaAlaValLeuPheile 858
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                                                                                                                                                                                                                                                                                             897 AsnValLeuSerProGlnGlnProSerArgPheCysSerLeuHisSerThrSerValCys
                                                                                                                                                    LysAlaGluGluThrLys-----GlnGlnIleAsnLysLeuAsnSerGluValThrThr
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917 ProSerArgGluSerLeuGlnThrArgThrSerTrpSerAlaHisGlnProCys-----
                                                                                                                                                                                                                                                                                                                                                                                                                                  membrane protein;
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drug screening; hypertension; renal failure; diabetes insipidus;
diabetic nephropathy; hypothyroidism; goiter; hypoparathyroidism;
pancreatic insufficiency; diabetes mellitus; cystic fibrosis;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          1059 ThrValSerSerPheSerLeuGluAsnLeuProGlySerTrpAsnGlnGlu 1075
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sialorrhea; salivary insufficiency; membrane potential; current flow; ion flux; transcription; signal transduction; assay; detection;
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       Novel mammalian potassium channel genes and polypeptides encoded by them for screening drugs useful for treating diseases such as hypertension, acute renal failure, diabetes insipidus and
                                                                                                                                                                                                                                                                                                                                                          (UYNY ) UNIV NEW YORK STATE RES FOUND.
                                                                                                                                                                                                                                                          99WO-US19902.
                                                                                                                                                                                                                                                                                                         98US-0098413.
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               hypothyroidism
                                                                                                    Rattus rattus.
                                                                                                                                                                                                                                                        31-AUG-1999;
                                                                                                                                                                                                                                                                                                         31-AUG-1998;
     sialorrhea;
                                                                                                                                                                                                      09-MAR-2000
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FX X X

Potassium channel genes e.g. elkl, elk2 or eag2 are useful for identifying modulators which are useful for treating hypertension, acute and chronic renal failure, diabetes insipidus, diabetic nephropathy, hyperthyroidism, goiter, hyperparathyroidism, pancreatic insufficiency, diabetes mellitus, cystic flancis, sialorrhea, salivary insufficiency. The availability of the gene sequences provides a tool for research into the physiobiological characteristics of the various genes and proteins for potassium channels including the development of medicines effective for treating disease conditions associated with mutations or defects in potassium channels and the screening of drugs to ensure that potassium channels are not blocked or physiologically affected by those drugs. The channel proteins encoded by these genes are also useful themselves as reporter molecules in assay and detection systems to measure changes in potassium concentration, membrane potential, current flow, ion flux, transcription, signal transduction, receptor-ligand interaction and second messenger

Claim 29; Page 82-86; 102pp; English.

1102 AA; Seguence

concentrations.

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1102
551
149
295
134
24
                              Conservative:
                                       Mismatches:
Indels:
                     Matches:
            1.53e-155
                   2515.00
62.00%
48.80%
41.30%
                                    Best Local Similarity:
                           Percent Similarity
Aliqnment Scores:
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US-09-965-830-1_COPY_6_3257 (1-3252) x AAY83028 (1-1102)

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ATGCCGGCCATGCGGGCCTCCTGGCGCCTCAGAACACCTTCCTGGACACCATCGCTACG 60		61 GGTTCGACGGCACGCACAGTAACTTCGTGGTGGCAACGCCCAGGTGGCGGGGCTCTTC 120	21 ArgPheAspGlyThrHisSerAsnPheIleLeuAlaAsnAlaGlnValAlaLysGlyPhe 40	121 CCCGTGGTCTACTGCTCTGATGGCTTCTGTGACCTCACGGGCTTCTCCCGGGCTGAGGTC 180	41 ProlleValTyrCysSerAspGlyPheCysGluLeuAlaGlyPheAlaArgThrGluVal 60
٦.	Т	61	21	121	41
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ATGCAGCGGGGCTGTGCCTCCTTTTATGGGCCAGACACCAGTGAGCTCGTCCGC 240

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80 300 100 360 120 ` 140 477	537 176 597 196 657 717 717 777	837 275 295 295 315 1017 1017	1137 375 1197 395 1257 1257 1317
MetGlnLysSerCysSerCysLysPheLeuPheGlyValGluThrAsnGluGlnLeuMet CAACAGATCGGCAAGGCCTGGACGACGACGACTCAAGGCTGACCTGATCTTL	TCCAAAGGCTTCAATGCCAACCGGCGGGCGGGCCGGGCC		CTGACACTGCTCATGGCCGTGCTCCCCCCCCCCCCCCCC
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TGGAGCTGCTGGCGGCCCGT	118 CTGGAGCTGCGGGGGGGGGGGGGGGGGGGGGGGGGGGGG
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Qy	GGCAGGGCAGGGGCTTTG	
đ	769 ValSerSerAlaAsnSerProLysThrLysGlnGluAlaAspProPro 784	
δλ	2395 GCTCCCCCACGGGCCCTAGAGGGGCTACGGCTGCCCCCATGCAT 2445	
qq	785 AsnHisGlyThrArgLysGluLysAsnLeuLysValGlnLeuCysSerLeuGlyThrAla 804	
Qy Db	2446 GIGCCCCCAGATCTGAGCCCCAGGGTAGTAGATGGCATTGAAGACGGCTGTGGCTCGGAC 2505	
ογ.	2506 CAGCCCAAGTTCTCTTTCCGCGTGGCCCAGTCTGGCCCCGGAATGTAGCAGCAGCCCCC	
8	825 GluThrGlnTh	
Oy Op	2563 TCCCTGGACCAGAGAGCGGCCTGCTCACTGTTCCCCATGGCCCCAGCGAG 2613	•
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7 g	### 1980 GluValSerGlnLeuGlyLysAspMetArgSerIleMetGlnLeuLeuGluAsnIleLeu 899	
ΟŊ	2731 GCGCCCACAGGGAGGTCCGTGCCTCGGGCATCGGGAGAGGGGCCGTGCCCAGC 2790	
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qq	955SerValAspProSerLeuValGlySerAsnProGlnArgThrGluAlaHisGlu 972	
Qγ	2950 AGCICCCCC 2958	
qq	973 GlnSerProValAspSerGluLeuHisHisSerProAsnLeuAlaTyrSerProSerHis 992	
δy	2959TGGCCTCGAGCCACA 2973	
đ	993 CysGlnValIleGlnGluGlyHisLeuGlnPheLeuArgCysIleSerProHisSerAsp 1012	
δ	4	
đ	3 ThrThrLeuThrProLeuGlnSerIleSerAlaThrLeuSerSerVerY	:
δλ	₹	
q	1033 SerGluThrSerLeuHisLeuValLeuProSerArgSerGluGluGlySerTleThrHis 1052	
οy	GCTGAGGCTACCAGCACTGGAGAGCCCCCAC	
đ	1053 GlyProValSerSerPheSerLeuGluAsnLeuProGly 1065	
ΟŅ	-	
a	Ser	
RESU AAY2 ID	RESULT 12 AAY22427 UD AAY22427 standard; Protein; 1017 AA.	
AC A	AAY22427;	· ·

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Brain specific potassium channel; human; central nervous system disorder; dementia; cerebral ischaemic sclerosis; therapy.
                                                                                                                                                                                                                                                                                                                                                                                                                                    This sequence is the potassium channel protein of the invention, that is expressed specifically in brain tissue. The protein is used to treat and investigate disorders of the central nervous system such as dementia and cerebral ischaemic sclerosis.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   121 CCCGTGGTCTACTGCTCTGATGGCTTCTGTGACCTCACGGGCTTCTCCCGGGCTGAGGTC 180
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                                                                                                                                                                                                                                                                                                                                                                   Potassium channel protein expressed specifically in brain tissue and method for its production
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            1 ATGCCGGCCATGCGGGGCCTCCTGGCGCCTCAGAACACCTTCCTGGACACCATCGCTACG 60
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                                     Human brain specific potassium channel protein sequence.
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Mismatches:
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Matches:
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                                                                                                                                                                                                                                                                   (YAMA ) YAMANOUCHI PHARM CO LTD.
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            28-SEP-1999 (first entry)
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N-PSDB; AAX84911.
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                                                                                                            Homo sapiens
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23-JAN-1998;
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QQ	395 I	::: LeuTrpAspileGlyTrpLeuHisGluLeuGlyLysArgLeuGluValProTyr 41	7
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<u>ئ</u> و	1297 4	AGCGAGGCCAACGGGACGGGGCTGGAGCTGCTGGGGGGCCCGTCGTGCGGCGCGAGCGCCTAC 1:	356
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6 6 7	1357 2	ATCACCTCCCTCTACTTCGCACTCAGCACCTCACCAGCGTGGGCTTCGGCAACGTGTCC 14 :::	416 46
à	1417 9	GCCAACAGGACACGGAGAAGAICTICICCATCIGCACCAIGCTCAICGGGGGCCCTGAIG	476

qq	47 AlaAsnThrAspAlaGluLysIlePheSerIleCysThrMetLeuIleGlyAlaLeuMet 466
	1477 CACGCGGTGGTGTTGGGAACGTGACGGCCATCATCCAGCGCATGTACGCCCGCC
	37 CIGIACCACACGCACGCGCACCTGCGCGACTACATCCGCATCCGCGTATCCCCA
	CCCTCAAGCAGCGCATGCTGGAGTACTTCCAGGCCACTGGGGCGCTGAACAATGCCATC 16
	507 ProceutysGlnArgMetLeuGluTyrPheGlnThrThrTrpAlaValAsnSerGly11e 526
	1657 GACACCACCGAGCTGCTGCAGAGCCTCCCTGACGACGTGCGCGCACACATCGCCATGCAC 1716
	1717 CTGCACAAGGAGGTCCTGCAGCTGCCACTGTTTGAGGCGGCCAGCCGCGGCTGCCTGC
	77 GCACTGTCTCTGGCCCTGCGGCTTCTGCACGCCGGCGGCGACTACCTCATCCACCAA 18
	37 GGCGATGCCCTGCAGGCCCTCTACTTTGTCTGCTCTGGCTCCATGGAGGTGCTCAAGGGT 1
	7 GGCACCGTGCTCGCCATCCTAGGGAAGGCGACCTGATCGGCTGTGAGCTGCCCGGCGG 19
	607 AsnMetValLeuAlalleLeuGlyLysGlyAspLeuIleGlyAlaAspIleProGluPro 626
	1957 GAGCAG
	1987 AAGGGGCTGACGTACCTGCAGTGTCTGCAGCTGGCCTGCACGACGACCTT 2046
	2047 GCGCTGTACCCCGAGTTTGCCCCGCGCTTCAGTCGTGGCCCTCCGAGGGGAGCTCAGCTAC 2106
	2107 AACCTGGGTGCTGGGGGGGGCTCTGCAGGGTGGACACCAGCTCCTG 2154
	55AGCGGCGACAATACC. 2
	702 ArgSerProArgLeuSerGlnProArgSerGluSerLeuGlySerSerSerAspLysThr 721
	70 CITATGICCACGCIGGAGGAGAAGAACAGAIGGGGAGCACAGGGCCCCACGGICTCCCCA 22
	22 LeuproSerIleAlaGluPro 734
	2230 GCCCCAGCTGATGAGCCCTCCAGCCCCTGCTGCTGCTGCTGCTGCTCTCTCA 2289 2230 GCCCCAGCTGATGAGCCTCCAGCCCCTGCTGCTTGCTTGC
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2413 GAGGGG	2575 GAGAGCGCCTGCTACTGTTCCCCATGGGCCCAGCGAGGAACACAGACACA :::	2 CTGTGTGTGGACCTCCTCTGCTCCGGCTTCTGCGGCTT 2 CTGTGTGTGGACATCTCCTACTGCTGCTGCACCCCCAGCTGCTTG 2 CTGTGTGTGGACATCTCTCTACTGCTGCTGCAGCTGGCTTGTTTG 1	928 oCysProGlnLeuArgProProCysLeuSerProCysAlaSerArgProProProProCysLeuSerProCysAlaSerArgProProProProProCysLeuSerProCysAlaSerArgProProProProCysLeuSerProCysAlaSerArgProProProProProCysAlaSerArgProProProProCysAlaSerArgProProProCysAlaSerArgProProProCysAlaSerArgProProCysCorpaceArgCorp	LT 13 AAY77739 standard; Protein; 457 AAY77739; 19-MAY-2000 (first entry) Human ESK1 (hESK1) protein frag ESK1; eag similar K+ channel; p neurological; Alzheimer's disea	der; schlzophrenia; Huntington's musculoskeletal; proliferative; c opic; neuroprotective; antidepres antiparkinsonian; cardiant; cyto
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This is an eag similar K+ channel (ESK) polypeptide (hESK1) fragment. The hESK1 protein can be expressed by standard recombinant methodology. The ESK polypeptide, polynucleotides and antibodies are useful for treating and diagnosing various potassium channel associated disorders such as neurological disorders, e.g. Alzheimer's disease, depression, anxiety, paric, obsessive-compulsive disorders, attention deficit, epilepsy; hyperactivity disorders, autism, schizophrenia, Huntington's disease and Parkinson's disease, cardiovascular disorders, musculoskeletal disorders and proliferative disorders such as cancer. The ESK polynucleotide is also useful for synthesis of ESK and gene mapping. The polypeptide can be antibodies, peptides or other molecules such as synthetic drugs, autibodies, peptides or other molecules which have an effect on the
                                                                                                                                                                                                                                                                                                                                                                                                Novel eag similar potassium channel polypeptide useful for treating various neurological, cardiovascular, musculoskeletal and proliferative
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                                                             13-AUG-1999;
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New isolated nucleic acid detection reagent for detecting 1000 or more genes from Drosophila and for elucidating cell signalling and cell-cell
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                                                                                                                                                                                                                                                                                                                                                                                                                         insecticides, therapeutics and pharmaceutical drugs. The invention discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA sequences (ABL01840-ABL16175) and the encoded proteins
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                      Drosophila; developmental biology; cell signalling; insecticide; pharmaceutical.
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Drosophila melanogaster polypeptide SEQ ID NO 10494
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QY Db	1342	AGCAGCCTCACCAGCGTGGGC 14 :::
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Qy Dp	1462	TGACGCCCATCATCCAGCGCATG 15:
Qy Db	1522 539	CTGTACCACAGCCGCGCGCCTGCGCGACTACATCCGCATC 15
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Oy Db	1942 (UValAlaThrSerAsnGlvGlnMe+mhral=mhrmh-as-co-
Oy Op	1948 C	GCAGGTGGTAAAGGCCAATGCCGACGTGAAGGGCT
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seent sequence is a partial ERG-like protein 2
22) which is a member of ERG potassium channel family. This
22 is a result of initial sequencing of clone jlhbaa042h05 which
22 is a from human brain library. ERG-LP2 gene is mapped to
21.00monosome 3p21.3-24.3, between markers WI-4218 and RP_LI5_1. It
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303..327
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423..442
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oy Dp	121	CCCGTGGTCTACTGCTCTGATGGCTTCTGTGACCTCACGGGCTTCTCCCGGGCTCAGGTC 180	
0 <u>y</u> Db	, 181	ATGCAGCGGGGCTGTGCTTCCTTTATGGGCCAGACACCAGTGAGCTCGTCGC 240	
Q D	241	CAACAGATCCGCAAGGCCCTGGACGAGCACAAGGAGTTCAAGGCTGAGCTGATCCTGTAC 300	
QQ QD	301	GGGAAGAGGGGCTCCCGTTCTGGTGTCTCCTGGATGTGATACCCATAAAGAATGAGAA 36 ::: :::	
Qy Dp	361	GGGGAGGTGGCTCTTCCTAGTCTCTCACAAGGACATCAGCGAAACCAAGAACCG :: ::: :: GlyaspValValLeuPheLeuAlaSerPheArgAspIleThrAspThrLysValLy	
Qy Dp	421	GAGACAGGTGGTGGCGGCGCGATATGGCCGGGCA 	
Q P	481	CCACC : rHisi	
QY Up	538	CACCTGCAGAAGCAGCCAAGGGCAAGCACAAGCTCAATAAGGGGGTGTTTGGGGAGAAA 597 	
Qy Db	598	CCAAACTIGCCTGAGTACAAAGTAGCGGCATCCGGAAGTCGCCCTTCATCCTGTTGCAC 657 	
Q. Q	658	TGTGGGGCACTGAGAGCCACCTGGGATGGCTTCATCCTGCTCGCCACACTCTATGTGGCT 717	
Oy Dp	718	GTCACTGTGCCCTACAGCCTGTGTGTGAGCACAGCAGGGAGCCCAGTGCCGCGCGC 777 	
Qy Db	778 257	CCGCCCAGCGTCTGTGACCTGGCCGTGGAGGTCCTTCATCCTTGACATTGTGCTGAAT 837 ::: :::	
oy Db	838	TTCCGTACCACATTCGTGTCCAAGTCGGGCCAGGTGGTGTTTGCCCCAAAGTCCATTTGC 897 	
Qy Db	898		٠.
oy Ob	958	IACTTCGGGGCCCATCTGCTGAAGACGGTGCGCCTG 1	
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Search completed: May 7, 2003, 15:21:08 Job time: 230.5 secs

::: ::::: pSerAsnGluValMet 527	GluTyrPheGlnThrThrTrpSerValAsnAsnGlyIleAspSerAsnGluValMet	509	Ωp
CACCACCGAGCTGCTG 1674	GAGTACITCCAGGCCACCTGGGCGGTGAAACGATGGCATCGACACCACCGAGCTGCTG	1618	δλ
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CCTCAAGCAGCGCATGCTG 1617	GACCIGCGCGACIACAICCGCAICCACGIAICCCCAAGCCCCICAAGCAGCGCCAIGCIG	1558	δŏ
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GTACCACAGCGCACGCGC 1557	GTGACGCCATCATCCAGCGCATGTACGCCCGCCGCTTTCTGTACCACAGCCGCACGCGC	1498	δ
sAlaLeuValPheGlyAsn 468		449	qq
CGCGGTGGTGTTGGGAAC 1497	ATCTTCTCCATCTGCACCATGCTCATCGGCGCCCTGATGCACGCGGTGGTTTGGGAAC	1438	δλ
	LeuSerLeuThrSerValGlyPheGlyAsnValSerAlaAsnThrAspAlaGluLys	429	đ
CAACACGGACACCGAGAAG 1437	CTCAGCAGCCTCACCAGCGTGGGCTTCGGCAACGTGTCCGCCAAACACGGACACGAGAAG	1378	οy
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411	GlyAsnAsnThr	408	QO
CGAGGCCAACGGGACGGGG 1317	GGGAACAGCTCCGGCCAGAGTGACAACTGCAGCAGCAGCAGCGAGGCCAACGGGACGGG	1258	δŏ
407	HisGluLeuGly	396	qq
GGGCCGGAGGCCAGCTGGA 1257	1198 CAGGAGCTGGCCCGCCGACTGCAGACTCCCTACTACCTGGTGGGCCGGAGGCCAGCTGGA 1257	1198	δ

